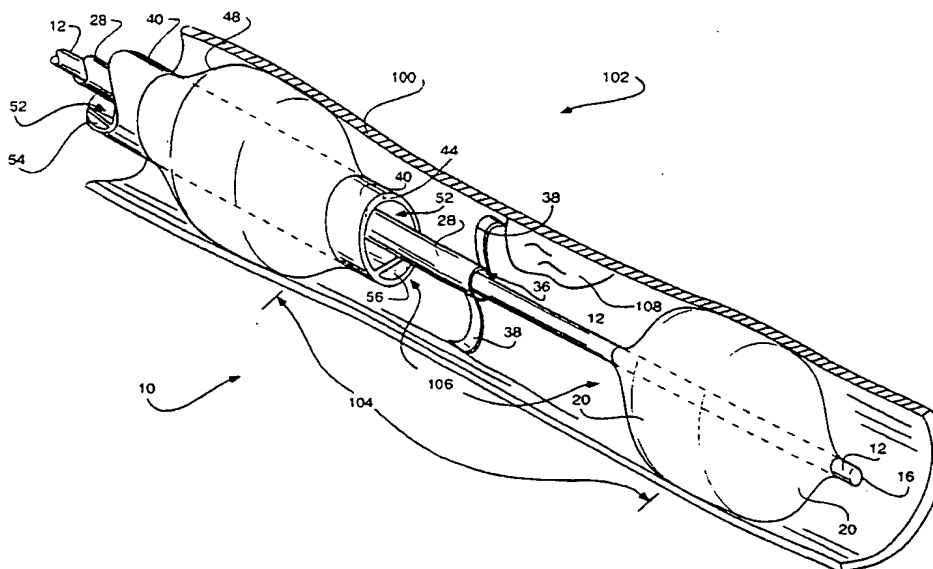


## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(54) Title: ADJUSTABLE TREATMENT CHAMBER CATHETER



## (57) Abstract

Disclosed herein is a catheter device (10) for treatment of disease in biological conduits. The device has inner (12) and outer (40) coaxial catheter members, each having an inflatable balloon (20, 48) attached near its distal end. When the balloons are inflated, a sealed treatment chamber (106) is created between the balloons. The length of the treatment chamber (106) is adjustable by sliding the coaxial catheter members (12, 40) with respect to each other to obtain a desired separation (104) between the attached balloons (20, 48) before inflation. Biological debris is trapped within the chamber (106) and removed by infusion and aspiration of a flushing fluid, reducing the risk of myocardial infarction. Adjunctive treatment devices can be inserted into the sealed treatment chamber through a lumen (52) of the outer coaxial member (40).

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## ADJUSTABLE TREATMENT CHAMBER CATHETER

### Technical Field:

This invention relates to catheters and more particularly to multiple concentric catheters having expandable balloons defining a treatment chamber.

### Background Art:

Blood vessels at certain critical parts of the human circulatory system can and often do become burdened with deposits, coatings, occlusions, and the like (hereafter "burden") which can reduce the vessel's blood carrying capacity. In some patients, this reduced capacity can cause serious and permanent injury. As a result, when significant burden is found to exist, some form of medical intervention is often indicated.

A number of intervention techniques have been developed over the years. These techniques permit the removal or reduction of the burden so that an increased and healthier blood flow can result. When a normal flow cannot be restored by these methods, the burdened portions of the vessel are sometimes bypassed using a vessel taken from elsewhere in the body. When the original blood vessel is a coronary artery, the bypass procedure is referred to as a coronary artery bypass graft.

20 The bypass graft itself may in time become burdened, as was the original vessel, therefore requiring further intervention. One type of burden that is of particular concern in treatment of a bypass arises when a vein has been used to make the bypass graft. Venous grafts are particularly susceptible to deposits of a friable material which is easily loosened during an intervention.

25 The friable material is soft and crumbly, and pieces are easily dislodged from the inside of the bypass graft. Often a treatment site will include a high proportion of such friable material surrounding other types of burden. Under these circumstances, attempts to remove the other burden often result in much of the friable material breaking loose and entering the blood stream. This loosened biological debris can move downstream, completely blocking another portion of the artery and causing myocardial infarction.

The intervention techniques referred to above include use of known devices such as balloon angioplasty, rotational atherectomy, the transluminal extraction

catheter, stents, directional coronary atherectomy, and the like. Each of these techniques, and others, are favored by practitioners in the treatment of diseased saphenous vein grafts and other vessels. Yet, each of these techniques exposes a patient to some greater or lesser degree of risk of embolism. Unfortunately, 5 this risk is present under the most carefully conducted intervention because the friable material can be so easily dislodged.

The friable material is especially vulnerable to being dislodged from the vessel walls when a catheter is advanced across a region of diseased tissue.

Consequently, catheter type treatment devices which must be repeatedly 10 inserted and withdrawn are prone to damage or break loose such friable material.

The loosened material, if not trapped in some way, is then carried away in the blood stream.

In treating diseased vessels, it will be desirable to stabilize a treatment site before proceeding with other forms of intervention. The treatment site is 15 "stabilized" by removing as much of the friable material as possible. No current or known device satisfactorily addresses this problem.

Additionally, it is sometimes desirable to administer a medication at a localized treatment site within a vessel. However, the natural biological fluids of the vessel both dilute the medication and carry it away from the region of application.

20 What is needed is a device for stabilizing a treatment site by removing friable material before other intervention is begun.

Additionally, what is needed is a device for preventing loosened particles of friable material or other plaque burden from entering the blood stream.

There is a need for a device which minimizes the necessity of repeatedly 25 passing a device through a treatment site during the course of an intervention.

There is an additional need for a device which permits a localized application of a medication at a treatment site without undue dilution or the risk of the medication entering the blood stream.

**Disclosure of Invention:**

30 It is an object of this invention to provide a catheter device having a fluid sealed treatment chamber, including infusion and aspiration, which can be used to stabilize a treatment site.

It is an additional object of this invention to provide such a treatment chamber catheter device which remains in place at the stabilized treatment site, and which allows access by adjunctive intervention devices.

It is yet an additional object of this invention to provide a treatment chamber  
5 catheter device which allows infusion of medications, confines these to the treatment site and prevents their dilution.

In accordance with the above objects and those that will be mentioned and will become apparent below, the adjustable treatment chamber catheter of the present invention comprises:

10 inner and outer coaxial catheter members, both members having a distal end and a proximal end and being slidable with respect to each other;

two expandable sealing members, one sealing member being disposed near the distal end of each catheter member; and

means for expanding the sealing members to form a pair of seals with the  
15 biological conduit, an adjustable length treatment chamber being defined within the conduit and between the seals,

whereby, the adjustable treatment chamber catheter may be inserted into the biological conduit and the catheter members may be slid with respect to each other to extend the distal end of the inner catheter member beyond the distal end  
20 of the outer catheter member and to adjust the longitudinal separation between the sealing members to a predetermined distance, the sealing members then being expanded, the adjustable length treatment chamber is created within the conduit and between the seals.

In a preferred embodiment, the adjustable treatment chamber catheter  
25 according to the present invention further includes:

an intermediate coaxial catheter member located between the inner and the outer coaxial catheter members and having a distal end and a proximal end, the intermediate catheter member being slidable and rotatable with respect to the other catheter members; and

30 burden loosening means disposed near the distal end of the intermediate catheter member for sliding and axial rotation within the treatment chamber to loosen plaque burden deposited on the biological conduit.

In another preferred embodiment, the expandable sealing members define proximal and distal ends of the treatment chamber and the device includes infusion and aspiration for flushing the treatment site and for removing biological debris from the treatment chamber.

5 In another embodiment of the present invention, a treatment chamber catheter for insertion into a biological conduit comprises:

a first catheter member having proximal and distal ends and an outer surface, an inflatable sealing member being disposed on the outer surface at the distal end, and an inflation lumen having an opening for communication with the  
10 sealing member and extending from the proximal end to the opening;

a second catheter member having proximal and distal ends and an outer surface, a burden loosening member being disposed on the outer surface at the distal end, and an infusion lumen extending from the proximal to the distal end and loosely enclosing the first catheter member proximally of the distal sealing  
15 member and being adapted for axial rotation about the first catheter member; and

a third catheter member having proximal and distal ends and an outer surface, a proximal inflatable sealing member being disposed on the outer surface, an inflation lumen having an opening to the proximal inflatable sealing member and  
20 extending from the proximal end to the opening, and an aspiration lumen extending from the proximal end to the distal end and loosely enclosing the second catheter member proximally of the burden loosening members, a treatment chamber being defined within the conduit and between the expandable sealing members.

25 In a preferred embodiment, the diameter of the infusion lumen is larger than the diameter of the enclosed first catheter member. This difference in diameter allows a fluid to be passed through the infusion lumen and into the treatment chamber. The difference in diameter also permits the second catheter to be rotated at high speed about the first catheter member so that the burden

30 loosening members can be used to remove friable tissue surrounding a lesion.

The second catheter lumen is connected to an infusion port at its proximal end.

The port is used to introduce a flushing fluid or liquid medication or the like into the treatment chamber.

In another preferred embodiment, the diameter of the aspiration lumen is substantially greater than the diameter of the enclosed second catheter member.

- 5 The aspiration lumen is used for access by other catheter type devices to the treatment chamber and to extract excess fluid and biological debris. The infusion and aspiration lumina are connected to infusion and aspiration ports at their proximal ends.

It is an advantage of this invention to provide a catheter device having a  
10 treatment chamber whose length can be easily adjusted to seal a specific region of a blood vessel only.

It is an additional advantage of this invention to trap loosened particles and thereby to reduce a risk of embolism.

It is also an advantage of this invention to stabilize a treatment site in  
15 saphenous vein graft repair.

It is yet another advantage of this invention to confine an infused medication to a treatment site and to prevent dilution or the carrying away into the blood stream of the medication.

It is an additional advantage of this invention to provide an adequately flushed  
20 and aspirated treatment chamber.

It is an advantage of this invention to remain in place while other interventions use the treatment chamber.

**Brief Description of Drawings:**

For a further understanding of the objects and advantages of the present  
25 invention, reference should be had to the following detailed description, taken in conjunction with the accompanying drawing, in which like parts are given like reference numerals and wherein:

Fig. 1 is a partial side view of the adjustable treatment chamber catheter according to the present invention.

30 Figs. 2-6 are cross-sectional views taken through Fig. 1 in the direction of the arrows.

Fig. 7 is a partial perspective oblique view of the treatment chamber catheter illustrated in Fig. 1 deployed inside a blood vessel.

Fig. 8 is a partial side view of the adjustable treatment chamber catheter including an infusion lumen and an infusion opening near the distal end of the treatment chamber.

Fig. 9 is a partial side view of the apparatus illustrated in Fig. 8 showing coaxial catheter members being withdrawn into an outer catheter member.

Fig. 10 is a partial side view of the apparatus shown in Fig. 8 and including a guide wire.

#### 10 **Best Mode for Carrying Out the Invention:**

The words "proximal" and "distal," as used below, have the following meaning, the "proximal" end of the catheter device is the end farthest from the treatment chamber, while the "distal" end is the end closest to the treatment chamber and includes the parts illustrated in the drawing.

15 The phrases "blood vessel" and "biological conduit" are used interchangeably throughout. Though typically the treatment chamber catheter 10 will be employed within a blood vessel, it is not to be so limited. In general it can be used within any biological vessel or conduit of a body.

The invention will now be described with respect to Fig. 1, which illustrates the adjustable treatment chamber catheter, identified generally by the numeral 10, in accordance with the present invention. The adjustable treatment chamber catheter 10 includes concentric first, second and third catheter members, 12, 28 and 40, respectively, a distal expandable sealing member 20, a proximal expandable sealing member 48, and burden loosening members 38.

25 As illustrated in Fig. 1, each of the three concentric catheter members 12, 28 and 40, extends progressively beyond a distal end of the enclosing catheter member. The proximal expandable sealing member 48 is attached near the distal end of the third catheter member 40. The burden loosening members 38 are attached to the second catheter member 28 near its distal end. The distal expandable sealing member 20 is attached near the distal end of the first catheter member 12. The first and third catheter members, 12, 40, may be adjusted axially with respect to one another to adjust the separation between the



distal and proximal expandable sealing members, 20, 48. The expandable sealing members will be inflated to form fluid seals with the walls of a biological conduit. The second catheter member 28 and attached burden loosening members 38 will be rotated at high speed allowing the burden loosening members to loosen and remove diseased tissue. The loosened and removed tissue will be confined between the inflated sealing members, 20, 48.

The adjustable treatment chamber catheter 10 includes a first catheter member 12 having a proximal end 14 and a distal end 16. The first catheter member 12 has an outer surface 18 and a distal expandable sealing member 20 which is attached circumferentially on the outer surface 18 near the distal end 16. The distal expandable sealing member 20 has an interior 22. The first catheter member 12 also includes a first lumen 24 extending between the proximal and distal ends, the first lumen 24 having an opening 26 to the interior 22 of the distal expandable sealing member 20 for inflation of the member 20. The first lumen 24 terminates in a closure (shown generally as the distal end 16) distally of the opening 26.

A second catheter member 28 has a proximal end 30, a distal end 32, and an outer surface 34. The second catheter member 28 has a second lumen 36 extending between the proximal end 30 and the distal end 32 and loosely enclosing the first catheter member 12 from a point near its proximal end 14 to a point proximal to the distal expandable sealing member 20. The second catheter member 28 includes a plurality of burden loosening members 38 which are radially disposed on the outer surface 34 near the distal end 32.

A third catheter member 40 has a proximal end 42, a distal end 44, and an outer surface 46. The third catheter member 40 includes a proximal expandable sealing member 48 which is attached circumferentially to the outer surface 46 near the distal end 44. The proximal expandable sealing member 48 has an interior 50. A third lumen 52 extends between the proximal and distal ends of the third catheter member 40 and has a diameter which is substantially larger than the diameter of the enclosed second catheter member 28. The third lumen 52 encloses the second catheter member 28 from a point near the proximal end 30 to a point proximal to the burden loosening members 38. A fourth lumen 54

extends between the ends of the third catheter member 40 and has an opening 58 to the interior 50 of the proximal expandable sealing member 48 for inflation of the member 48. The fourth lumen 54 terminates in a closure 56 distally of the opening 58.

5 Figures 2-6 are cross-sectional views taken at corresponding lines through the partial catheter shown in Fig. 1 looking in the direction of the arrows.

With particular reference to Fig. 2, the concentric catheter members are shown to have a generally circular cross-section. The first catheter member 12 includes the first lumen 24. The second lumen 36 of the second catheter member 28 is 10 slightly greater in diameter than the first catheter member 12, which it encloses. The third lumen 52 of the third catheter member 40 is substantially greater in diameter than the second catheter member 28, which it encloses. A fourth lumen 54 is shown.

With particular reference to Fig. 3, there is illustrated a cross-section through 15 the proximal expandable sealing member 48 and the inflation opening 58 of Fig.

1. The interior 50 of the proximal expandable sealing member 48 communicates with the fourth lumen 54 via the opening 58.

With particular reference to Fig. 4, there is illustrated a cross-section taken at the line 4-4 of Fig. 1 and illustrates two burden loosening members 38 radially 20 attached to the outer surface 34 of the second catheter member 28. The second catheter member 28 is no longer enclosed by the third lumen 52 of the third catheter member 40, and is shown extending distally of the distal end 44 of the third catheter member 40.

With particular reference to Fig. 5, there is illustrated a cross-section through 25 the first catheter member 12 of Fig. 1 showing the first lumen 24. The second lumen 36 of the second catheter member 28 no longer encloses the first catheter member 12.

With particular reference to Fig. 6, this is an illustration of a cross-section taken through the distal expandable sealing member 20 and the inflation opening 26 of 30 Fig. 1. The interior 22 of the distal expandable sealing member 20 communicates with the first lumen 24 via the opening 26.

### In Use

The adjustable treatment chamber catheter 10 is indicated for the treatment of saphenous vein grafts where the removal of friable material within a sealed treatment chamber can be accomplished with minimum risk of embolization.

- 5 Once the treatment site has been stabilized, other forms of treatment can be used in conjunction with the present invention.

The present invention will now be described with reference to a typical intervention. A guiding catheter (not shown) will be introduced first via a femoral artery into the aorta. A guiding catheter has a single large lumen used for  
10 insertion of other catheter devices. The guiding catheter will then be left in place throughout the remainder of the intervention and will provide a safe and rapid access to the aorta region.

Next, the adjustable treatment chamber catheter 10 will be inserted through the guiding catheter and into a selected coronary bypass graft or artery. Fluoroscopy  
15 is typically used to guide the distal end of the catheter 10 to the desired treatment site.

When the situation at the treatment site permits, the location of the expandable sealing members 20, 48 will be adjusted to establish a treatment chamber of the desired length, with one sealing member located distally of the site and the other  
20 located proximally of the site, and a desired distance apart. The expandable sealing members 20, 48 will be inflated so that they form a fluid seal with the walls of the vessel and create the treatment chamber around the site.

Fig. 7 illustrates the treatment chamber 106 established at a treatment site 102 inside a vessel 100. The expandable sealing members 20, 48 are shown in the  
25 inflated condition, forming fluid seals with the walls of the vessel 100. Friable material 108 is shown attached to a wall of the vessel 100 at the treatment site 102. To loosen or remove the friable material 108, the second catheter member 28 is rotated at a high speed, and its longitudinal position is adjusted to bring the burden loosening members into contact with the material 108.

- 30 Infusion of a flushing fluid is made through the second lumen 36 from an infusion port (not shown) located at the proximal end 30 of the second catheter member 28. Loosened particles of friable material and excess fluid are removed

from the treatment chamber 106 back through the third lumen 52 and an aspiration port (not shown).

The diameter of the second lumen 36 exceeds the diameter of the first catheter member 12 just sufficiently to permit infusion to occur at a useful rate, and for the second catheter member 28 to rotate freely about the first catheter member 12.

The diameter of the third lumen 52 substantially exceeds the diameter of the second catheter member 28 so that aspiration of particles of loosened friable material and excess fluid can occur at a rate corresponding to the infusion rate. In general, the size of the particles will permit aspiration in the manner described.

10 When the size of the particles otherwise prevents removal by aspiration as described, the diameter of the third lumen 52 will permit insertion of another catheter device used to further reduce the size of the particles.

#### **Sealed Treatment Chamber**

The two inflated expandable sealing members 20, 48 prevent the release of loosened friable material from the treatment chamber 106 into the normal blood stream of the vessel 100. The risk that such particles might be carried away into the circulatory system and form emboli is greatly reduced, though not entirely eliminated. Such particles can still enter the blood stream and pose a risk whenever the treatment chamber catheter 10 is being positioned or repositioned.

20 This occurs during such times because the expandable sealing members are not inflated and therefore do not form a fluid seal with the walls of the vessel, and also because the device must be passed across a lesion at a new treatment site.

The existence of a sealed treatment chamber 106 having infusion and aspiration and an access lumen (the third lumen 52) of substantial diameter create a safe and flexible environment within which to perform additional intervention. Once the treatment site 102 has been stabilized by a removal of a majority of the unstable friable material 108, other burden can be addressed using preferred intervention techniques, such as balloon angioplasty, rotational atherectomy, the transluminal extraction catheter, stents, directional coronary

25 create a safe and flexible environment within which to perform additional intervention. Once the treatment site 102 has been stabilized by a removal of a majority of the unstable friable material 108, other burden can be addressed using preferred intervention techniques, such as balloon angioplasty, rotational atherectomy, the transluminal extraction catheter, stents, directional coronary

30 atherectomy, and the like. Each of these techniques, used in the absence of the sealed treatment chamber 106, encounters some difficulty in preventing the escape of particles into the blood stream. These are not only particles of

accidentally loosened friable material, as in a vein graft repair, but include particles of other types of burden removed as a primary object of the intervention.

Thus, the sealed treatment chamber 106 provides an accessible and continually flushed environment for adjunctive treatment of arterial heart disease.

- 5 The sealed treatment chamber substantially reduces the risk of embolism.

#### **Medications**

The sealed treatment chamber 106 also allows the introduction of specific medications by infusion to the treatment site 102. Such medications are confined at the treatment site 102. Normally, a medication in the form of a fluid could not  
10 be applied at a treatment site because there would be no practical way to prevent the medication from being carried away in the blood stream. The two expandable sealing members 20, 48 form fluid seals with the walls of a vessel 100, however, and thus effectively confine a medication to the treatment chamber 106. The medication can thus be applied in undiluted form directly at  
15 the treatment site 102. This ability to confine the application and to prevent dilution, greatly enhances the effectiveness of such treatment.

#### **Alternative Embodiments**

Fig. 8 illustrates a partial side view of an alternative embodiment of the treatment chamber catheter in accordance with the present invention. The  
20 arrangement for infusion of fluid into the treatment chamber provides flow from the distal end of the chamber toward the proximal end. This embodiment eliminates the need for an additional pressurized, rotating seal near the proximal end of the catheter device (not shown).

The alternative treatment chamber catheter 108 includes an outer coaxial  
25 catheter member 110, an inner coaxial catheter member 112, a pair of expandable sealing members 114, 116, an intermediate coaxial catheter member 118, and burden loosening members 120.

The outer coaxial catheter member 110 has a distal end 122 and a proximal end 124. The expandable sealing member 114 is disposed on the outer coaxial  
30 catheter member 110 near the distal end 122. The outer coaxial catheter member 110 includes an access and aspiration lumen 126, and an inflation lumen 128. The expandable sealing member 114 has an interior 130 for

receiving and storing a fluid during expansion. An opening 132 connects the inflation lumen 128 with the interior 130. The inflation lumen 128 extends from the proximal end 124 to the opening 132. During expansion of the expandable sealing member 114, a fluid is introduced at the proximal end of the inflation lumen 128. The fluid travels along the lumen 128, then through the opening 132 and into the interior 130, urging the expandable sealing member 114 against the walls of the biological conduit (not shown).

The inner coaxial catheter member 112 has a distal end 134 and a proximal end 136. The expandable sealing member 116 is disposed on the inner coaxial catheter member 112 near the distal end 134. The inner coaxial catheter member 112 includes an infusion lumen 138 and an inflation lumen 140. The expandable sealing member 116 has an interior 142 for receiving and storing a fluid during expansion. An opening 144 connects the inflation lumen 140 with the interior 142. The inflation lumen 140 extends from the proximal end 136 to the opening 144. The expandable sealing member 116 has a proximal end 146.

The inner coaxial catheter member 112 includes an infusion opening 148 located near the proximal end 146 of the sealing member 116. A fluid is introduced at the proximal end of the infusion lumen 138, travels along lumen 138, then through the infusion opening 148 (shown with arrow).

The expandable sealing member 114 has a distal end 150. A sealed treatment chamber 152 is defined between the expanded sealing members 114, 116. The access and aspiration lumen 126 extends from the proximal end 124 to the distal end 122 of the outer coaxial catheter member 110, opening into the treatment chamber 152. Excess fluid and biological debris are removed from the treatment chamber 152 through the access and aspiration lumen 126 (shown with arrow).

Fig. 9 is a partial side view of the apparatus illustrated in Fig. 8. The inner coaxial and the intermediate coaxial catheter members 112, 118 are shown slid proximally of the distal end 122 of the outer coaxial catheter member 110. The expandable sealing members 114, 116 are shown deflated, the inflation fluid having been withdrawn from interiors 130, 142. The burden loosening members 120 are shown folded within the distal end of the access and aspiration lumen

126. In this configuration, the catheter device 108 can be inserted, removed and repositioned within a biological conduit.

In another alternative embodiment, Fig. 10 is a partial side view of a treatment chamber catheter including a guide wire 156. The infusion lumen 138 includes a small opening 154 at the distal end 134. The guide wire 156 extends through the infusion lumen 138, then passes through the small opening 154, extending distally of the distal end 134. The guide wire 156 is slidable within the infusion lumen 138. The fit of the small opening 154 is adapted to allow the guide wire 156 to slide within the opening 154 while not allowing infusion fluid to pass beyond the distal end 134 and into the blood stream.

In another alternative embodiment, perfusion can be added by bypassing the treatment chamber with a perfusion lumen (not shown). This allows some blood to continue to flow within the vessel and around the sealed treatment chamber, reducing ischemic stress and permitting an intervention of longer duration.

15

#### **A Treatment Method**

The present invention readily lends itself to the treatment of atherosclerotic vessels. A method for conducting such treatment is defined which begins with a step of providing a treatment chamber catheter of the type described above.

The method continues with the steps of guiding the treatment chamber catheter into the biological conduit. Then determining a location of a treatment site. This step is usually accomplished by means of fluoroscopy and radio opaque dyes. Next, the distal sealing member is positioned distally to the selected treatment site, and the proximal sealing member is positioned proximally of the selected site. The separation of the two sealing members is adjusted by sliding the catheter members with respect to one another. Then, the sealing members are inflated to seal off the region of the conduit surrounding the site (see Fig. 7). Finally, a flushing fluid is passed through the sealed region by means of infusion and aspiration.

Having established a sealed, flushed treatment chamber, it is now desirable to operate the second catheter member 28 (Fig. 7) to cause the burden loosening members 38 to engage the friable material 108. It is desirable to remove as much of this material as possible from the site before beginning other types of

treatment. The loosened material will be trapped within the treatment chamber 106 and will then be removed by aspiration back through lumen 52.

Now that a stabilized treatment region has been prepared, another adjunctive treatment device can be inserted into the treatment chamber 106 (not shown).

- 5 As a preliminary step, a guiding catheter might be inserted between a point of entry into the body and a region near the treatment site. For example, the guiding catheter might be inserted between a point of entry at the groin and extend to the aorta (not shown).

While the foregoing detailed description has described several embodiments of  
10 the adjustable treatment chamber catheter device 10 in accordance with this invention, it is to be understood that the above description is illustrative only and not limiting of the disclosed invention. It will be appreciated that the specific arrangement of the burden loosening members can be different from that shown in the drawing and that the relative difference in diameter of the various lumina  
15 and catheter members can differ from the examples shown in the accompanying figures and remain within the scope and spirit of this invention. Also, it will be appreciated that the device can be used within any biological conduit of the body rather than being limited to use in a blood vessel. Thus the invention is to be limited only by the claims as set forth below.



**Claims:**

1. An adjustable treatment chamber catheter for insertion into a biological conduit, comprising:

inner and outer coaxial catheter members, both members having a distal end  
5 and a proximal end and being slidable with respect to each other;

two expandable sealing members, one sealing member being disposed near  
the distal end of each catheter member; and

means for expanding the sealing members to form a pair of seals with the  
biological conduit, an adjustable length treatment chamber being defined within  
10 the conduit and between the seals,

whereby, when the adjustable treatment chamber catheter is inserted into the  
biological conduit and the catheter members are slid with respect to each other to  
extend the distal end of the inner catheter member beyond the distal end of the  
outer catheter member and to adjust the longitudinal separation between the  
15 sealing members to a predetermined distance, the sealing members then being  
expanded, the adjustable length treatment chamber is created within the conduit  
and between the seals.

2. The adjustable treatment chamber catheter of Claim 1, further  
comprising:

20 an intermediate coaxial catheter member located between the inner and the  
outer coaxial catheter members and having a distal end and a proximal end, the  
intermediate catheter member being slidable and rotatable with respect to the  
other catheter members; and

burden loosening means disposed near the distal end of the intermediate  
25 catheter member for sliding and axial rotation within the treatment chamber to  
loosen plaque burden deposited on the biological conduit.

3. The adjustable treatment chamber catheter of Claim 1, further including  
infusion means for the infusion of a fluid into the treatment chamber, and  
aspiration means for the aspiration of a fluid and biological debris out of the  
30 treatment chamber.

4. The adjustable treatment chamber catheter of Claim 3, wherein the defined treatment chamber has a distal end, and the infusion means includes means for infusion of the fluid from the distal end of the treatment chamber.

5. The adjustable treatment chamber catheter of Claim 4, wherein the 5 means for infusion from the distal end includes the expandable sealing member disposed on the inner catheter member having a proximal end and a distal end, and the inner catheter member having an infusion opening near the proximal end of said sealing member, the inner catheter member further having an infusion lumen extending from the proximal end of the inner catheter member to the 10 infusion opening, whereby when a fluid is introduced at the proximal end of the infusion lumen, the fluid moves through the lumen toward the infusion opening, then through the opening and into the treatment chamber.

6. The adjustable treatment chamber catheter of Claim 3, wherein the defined treatment chamber has a proximal end and wherein the aspiration means 15 includes an aspiration opening near the proximal end of the treatment chamber for the aspiration of the fluid and the biological debris out of the treatment chamber.

7. The adjustable treatment chamber catheter of Claim 6, wherein the outer catheter member includes an aspiration lumen extending from the proximal end 20 of the outer catheter member to the aspiration opening into the treatment chamber, whereby when suction is applied at the proximal end of the aspiration lumen, fluid and biological debris are removed from the treatment chamber by passing through the aspiration opening then along the aspiration lumen toward the proximal end.

25 8. The adjustable treatment chamber catheter of Claim 1, further including guiding means for guiding the adjustable treatment chamber catheter within the biological conduit.

9. The adjustable treatment chamber catheter of Claim 8, wherein the guiding means includes the inner coaxial catheter member having an opening at 30 the distal end and a lumen extending from the proximal end to the opening, and a guide wire passing through the lumen and extending through the opening at

the distal end of the lumen, the guide wire being slidable within the lumen and through the opening.

10. The adjustable treatment chamber catheter of Claim 1, further including means for contracting the sealing members, whereby when the sealing members  
5 are contracted, the device can be more easily inserted, removed, and adjusted within the biological conduit.

11. The adjustable treatment chamber catheter of Claim 1, wherein the expandable sealing members each comprise an inflatable balloon, each inflatable balloon having an interior for receiving a fluid during expansion.

10 12. The adjustable treatment chamber catheter of Claim 11, wherein the means for expanding the sealing members includes the inner and the outer coaxial catheter members each having an inflation lumen and an opening connecting the lumen with the interior of the corresponding balloon, the inflation lumen extending from the proximal end of the catheter member to the opening,  
15 whereby when the catheter device is placed within a biological conduit and a fluid is introduced at the proximal end of the inflation lumen, it will pass through the lumen and then through the opening and into the balloon, thereby expanding the balloon, urging the balloon against, and forming a seal with, the biological conduit.

20 13. A treatment chamber catheter for insertion into a biological conduit, comprising:

a first catheter member having proximal and distal ends and an outer surface, an inflatable sealing member being disposed on the outer surface at the distal end, and an inflation lumen having an opening for communication with the  
25 sealing member and extending from the proximal end to the opening;

a second catheter member having proximal and distal ends and an outer surface, a burden loosening member being disposed on the outer surface at the distal end, and an infusion lumen extending from the proximal to the distal end and loosely enclosing the first catheter member proximally of the distal sealing  
30 member and being adapted for axial rotation about the first catheter member; and

a third catheter member having proximal and distal ends and an outer surface, a proximal inflatable sealing member being disposed on the outer surface, an inflation lumen having an opening to the proximal inflatable sealing member and extending from the proximal end to the opening, and an aspiration lumen  
5 extending from the proximal end to the distal end and loosely enclosing the second catheter member proximally of the burden loosening members, a treatment chamber being defined within the conduit and between the expandable sealing members.

14. The treatment chamber catheter of Claim 13, further including a  
10 separation being defined between the sealing members, the separation being adjustable by sliding the third catheter member with respect to the first catheter member, and an adjustable length treatment chamber being defined,  
whereby when the treatment chamber catheter is inserted into the biological conduit, the separation between the inflatable sealing members then being  
15 adjusted by sliding the first and the third catheter members with respect to each other, the sealing members then being inflated, an adjustable length treatment chamber is created within the conduit and between the seals.

15. The treatment chamber catheter of Claim 13, wherein the infusion lumen has a diameter and the first catheter member has a diameter and the diameter of  
20 the lumen exceeds the diameter of the catheter member by an amount sufficient to permit a fluid to pass through the lumen.

16. The treatment chamber catheter of Claim 13, wherein the aspiration lumen has a diameter and the second catheter member has a diameter and the diameter of the lumen substantially exceeds the diameter of the second catheter  
25 member.

17. The treatment chamber catheter of Claim 16 wherein the difference in diameters allows removal of fluid and biological debris from the treatment chamber through the aspiration lumen.

18. The treatment chamber catheter of Claim 13 wherein the second  
30 catheter member and the infusion lumen share a common longitudinal axis and wherein the second catheter member is adapted for rapid rotation about the first catheter member.

19. The treatment chamber catheter of Claim 15 wherein the second catheter member is adapted for rapid rotation about the first catheter member and the fluid passing through the infusion lumen.
20. A method for treating a diseased biological conduit, the method  
5 comprising the steps of:
- providing a treatment chamber catheter of the type having concentric catheter members and distal and proximal inflatable balloons for creating a sealed treatment chamber within a biological conduit and between the balloons, the length of the treatment chamber being adjustable by sliding the catheter  
10 members with respect to each other, the treatment chamber catheter including burden loosening means and both infusion and aspiration for removing biological debris and excess fluid from the treatment chamber, access to the treatment chamber by adjunctive catheter-type treatment devices being through a lumen;
  - guiding the treatment chamber catheter into the biological conduit;
  - 15 determining a location of a treatment site;
  - positioning the distal sealing member within the conduit distally to the determined treatment site;
  - positioning the proximal sealing member within the conduit proximally to the treatment site;
  - 20 adjusting the separation between the sealing members by sliding the concentric catheter members with respect to each other;
  - inflating the sealing members to create a sealed treatment chamber within the conduit and between the sealing members;
  - introducing a fluid into the chamber via the infusion lumen;
  - 25 removing excess fluid and biological debris from the chamber via the aspiration lumen.
21. The method of Claim 20 further including the step of inserting a guiding catheter from the groin into the aorta and continuing with the step of guiding the treatment chamber catheter through the guiding catheter and then into the  
30 biological conduit.
22. The method of Claim 20, further including the step of loosening burden deposited on the biological conduit.

23. The method of Claim 20 further including the steps of inserting an adjunctive catheter device for treatment through the aspiration lumen of the treatment chamber catheter and operating the inserted device within the sealed treatment chamber.

5 24. A fluid sealed treatment chamber catheter device for insertion into a biological conduit and having a treatment chamber of adjustable length, the catheter device comprising:

distal sealing means for creating a distal fluid seal within the biological conduit, the distal sealing means being attached near a distal end of a first catheter

10 member;

burden loosening means for the reduction of biological tissue at a treatment site, the burden loosening means being attached radially near a distal end of a second catheter member, the second catheter member including an infusion lumen, the lumen slidably and rotationally enclosing the first catheter member

15 proximally to the distal sealing means; and

proximal sealing means for creating a proximal fluid seal within the biological conduit, the proximal sealing means being attached near a distal end of a third catheter member, the third catheter member including an aspiration lumen having a diameter substantially greater than a diameter of the second catheter

20 member and enclosing the second catheter member proximally to the burden loosening means,

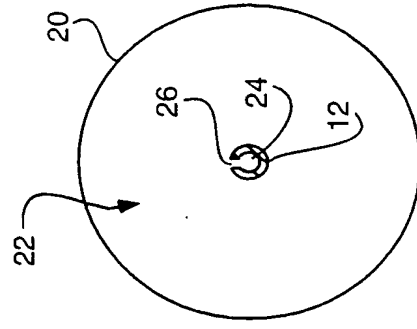
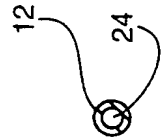
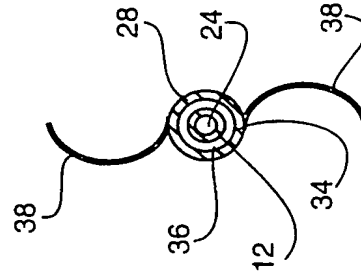
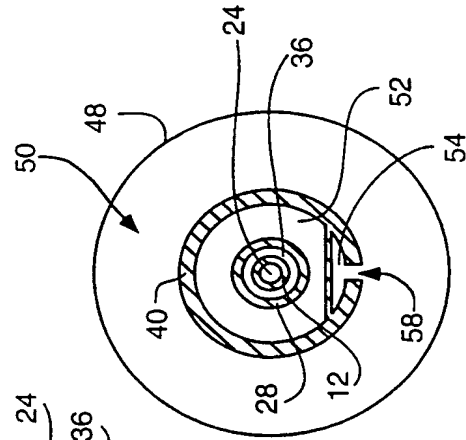
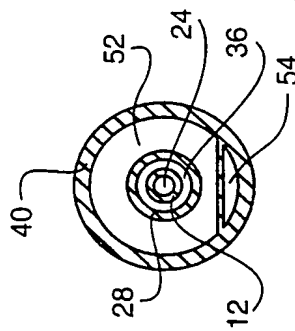
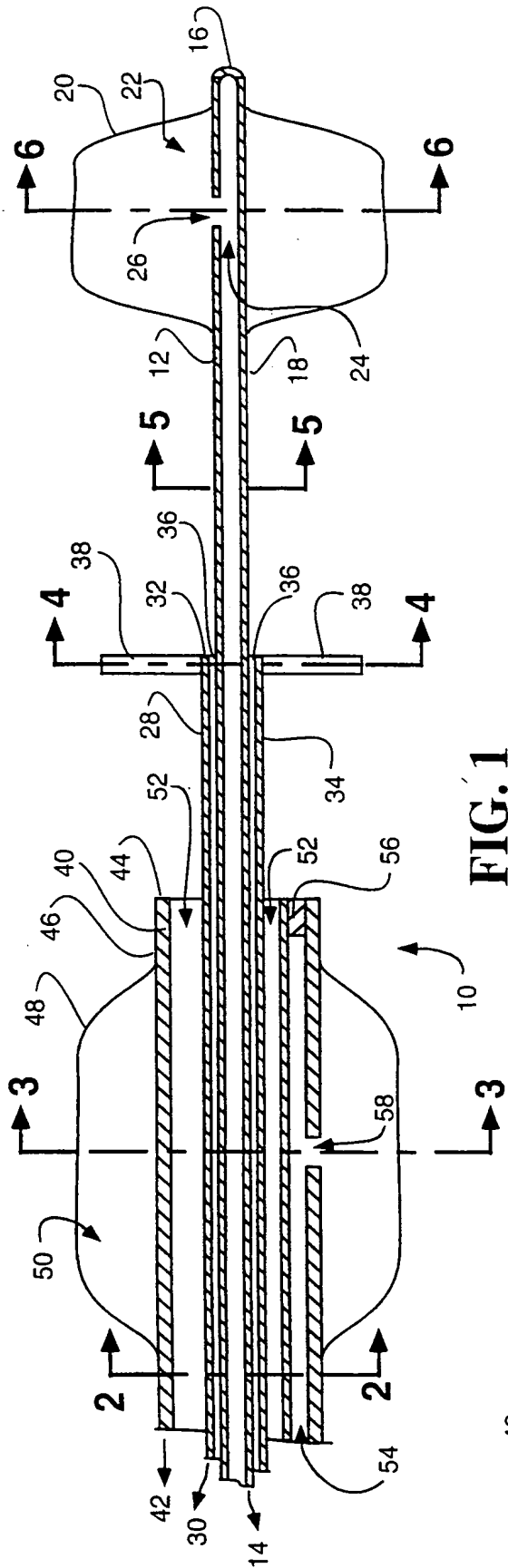
whereby, when the catheter device is inserted into a biological conduit, the distal sealing means being positioned distally to a treatment site, and the third catheter member being slid with respect to the first catheter member to locate the

25 proximal sealing means proximally to the treatment site and at an adjustable distance from the distal sealing means, a fluid sealed treatment chamber of adjustable length is thereby created within the conduit and between the sealing means.

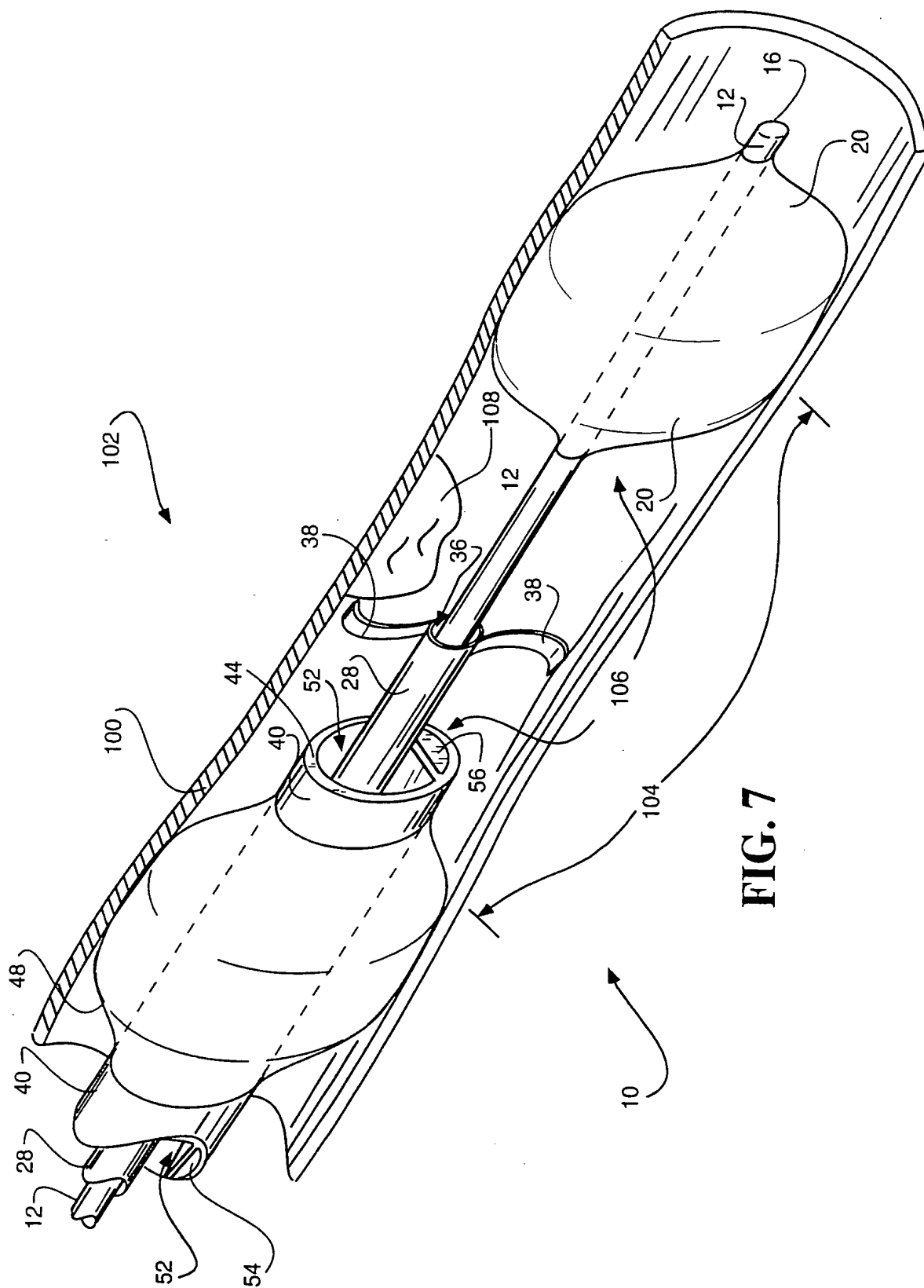
25. The catheter device of Claim 24 wherein the proximal and distal sealing

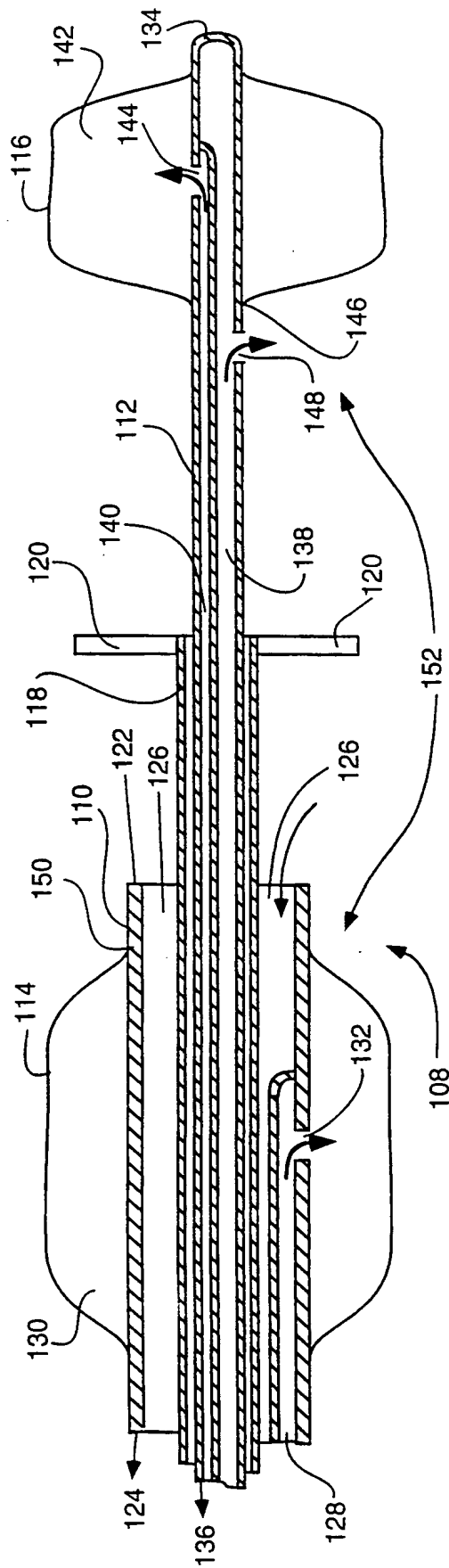
30 means are expandable balloons and the attached catheter members include inflation lumina for inflation of the balloons.

26. The catheter device of Claim 24 wherein the burden loosening means comprises a flexible member being disposed on the second catheter member.

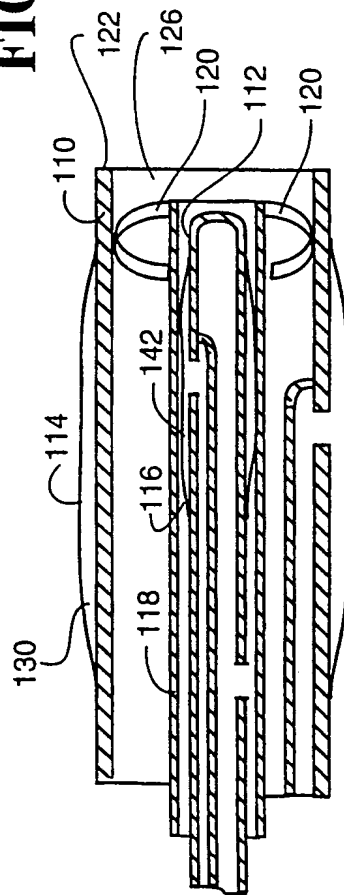




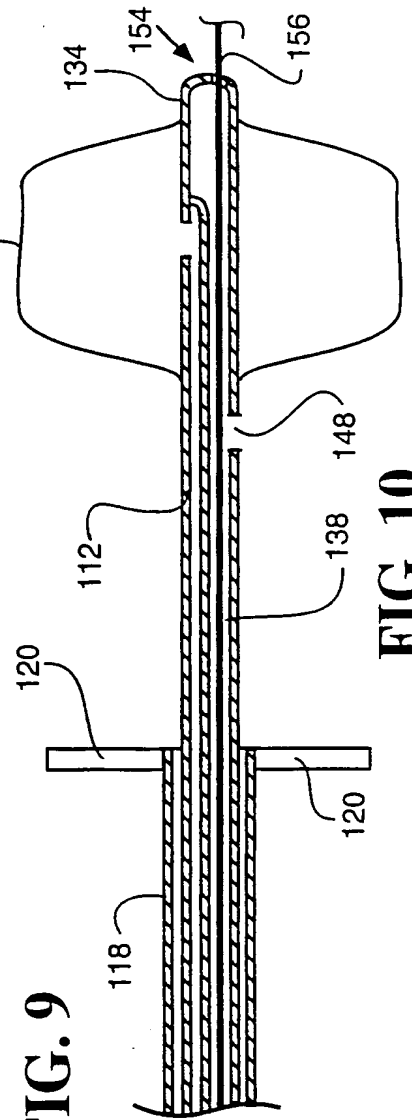




**FIG. 8**



**FIG. 9**



**FIG. 10**